



FORM PTO-1449 (REV. 7-80)	U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTY. DOCKET NO. 240083.501D4	EXPRESS MAIL NO. EL615486894US
		APPLICANTS Mary E. Brunkow et al.	
		FILING DATE October 25, 2000	GROUP ART UNIT

INFORMATION DISCLOSURE STATEMENT
(Use several sheets if necessary)

U.S. PATENT DOCUMENTS

*EXAMINER INITIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
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FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION	
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MB	AB	WO 98/45435	10/15/98	PCT				
	AC							
	AD							
	AE							
	AF							

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OTHER PRIOR ART (Including Author, Title, Date, Pertinent Pages, Etc.)

MB	AG	Barbosa et al., "Identification of the Homologous Beige and Chediak-Higashi Syndrome Genes," <i>Nature</i> 382:262-265, 1996.
↓	AH	Bignon and Siminovitch, "Identification of PTP1C Mutation as the Genetic Defect in Motheaten and Viable Motheaten Mice: A Step Toward Defining the Roles of Protein Tyrosine Phosphatases in the Regulation of Hemopoietic Cell Differentiation and Function," <i>Clinical Immunology and Immunopathology</i> 73(2):168-179, 1994.
	AI	Blair et al., "The Mouse Scurfy (sf) Mutation Is Tightly Linked to <i>Gata1</i> and <i>Tfe3</i> on the Proximal X Chromosome," <i>Mammalian Genome</i> 5:652-654, 1994.
	AJ	Database Genbank Accession No. AI949471, August 1, 1998.
	AK	Database Genbank Accession No. AJ005891, May 1, 1998.
	AL	Derry et al., "The Mouse Homolog of the Wiskott-Aldrich Syndrome Protein (WASP) Gene Is Highly Conserved and Maps near the Scurfy (sf) Mutation on the X Chromosome," <i>Genomics</i> 29:471-477, 1995.
↓	AM	Godfrey et al., "Fatal Lymphoreticular Disease in the Scurfy (sf) Mouse Requires T Cells that Mature in a sf Thymic Environment: Potential Model for Thymic Education," <i>Proc. Natl. Acad. Sci. USA</i> 88:5528-5532, 1991.
MB	AN	Godfrey et al., "X-Linked Lymphoreticular Disease in the Scurfy (sf) Mutant Mouse," <i>American Journal of Pathology</i> 138(6):1379-1387, 1991.

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							YES	NO
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M/S	BG	Lyon et al., "The Scurfy Mouse Mutant has Previously Unrecognized Hematological Abnormalities and Resembles Wiskott-Aldrich Syndrome," <i>Proc. Natl. Acad. Sci. USA</i> 87:2433-2437, 1990.
	BH	Rawlings et al., "Mutation of Unique Region of Bruton's Tyrosine Kinase in Immunodeficient XID Mice," <i>Science</i> 261:358-361, 1993.
	BI	Sugamura et al., "The Interleukin-2 Receptor γ Chain: Its Role in the Multiple Cytokine Receptor Complexes and T Cell Development in XSCID," <i>Ann. Rev. Immuno.</i> 14:179-205, 1996.
M/S	BJ	Veres et al., "The Molecular Basis of the Sparse Fur Mouse Mutation," <i>Science</i> 237:415-417, 1987.
	BK	
	BL	
	BM	
	BN	

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